IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant

CRIPPS et al.

Confirmation No: 1830

Appl. No.

10/774,393

Filed

February 10, 2004

Title

JAN 0 5 2005

PHARMACEUTICAL FORMULATION OF FLUTICASONE

PROPIONATE

TC/A.U.

: 1616

Examiner

: M Haghighatian

Docket No.:

: CRIP3001C3/REF

Customer No:

: 23364

TERMINAL DISCLAIMER

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Sir:

I, Richard E. Fichter, represent that I am the attorney of record for the aboveidentified application.

SmithKline Beecham Corporation (hereinafter referred to as "Assignee") is the owner of the entire right and title to the above-captioned patent application and the invention and improvements therein disclosed for the United States, by virtue of an assignment from the inventors to SmithKline Beecham Corporation dated March 31, 2001, and recorded on September 11, 2002. The assignee is also the owner of the entire right and title to U.S. Patent No. 6,479,035 issued on November 12, 2002 and U.S. Patent No. 6,630,129 issued on October 7, 2003, by virtue of the same assignment from the inventors to SmithKline Beecham Corporation. The present application and the '035 and '129 patents are based on the same initial filing in the United States.

The terminal part of any patent granted on the above-captioned application which would extend beyond the expiration date of the full statutory term of U.S. Patent Nos. 6,479,035 and 6,630,129 is hereby disclaimed. Any patent granted on the above-captioned application shall be enforceable only for and during such period that the legal title to said patent shall be the same as the legal title to the U.S. Patent Nos. 6,479,035 and 6,630,129, this agreement to run with any patent on the above-captioned application to be binding upon the grantee, its successors or assigns.

Appl. No. 10/774,393

Amendment dated: January 4, 2005 Reply to OA of: October 7, 2004

However, no rights available under the provisions of 35 U.S.C. §§155 and 156 are hereby waived.

No terminal part is disclaimed of any patent granted on the above-captioned application prior to the expiration date of the full statutory term of the U.S. Patent Nos. 6,479,035 and 6,630,129 in the event that it later expires for failure to pay a maintenance fee, is held unenforceable, is found invalid, is statutorily disclaimed in whole or terminally disclaimed under 37 C.F.R. § 1.321(a), has all claims canceled by a reexamination certificate, or is otherwise terminated prior to the expiration of its statutory term, except for the separation of legal title stated above.

The Terminal Disclaimer fee of \$130.00 is submitted herewith. Please charge any additional fees or credit any overpayments in connection with this paper to Deposit Account No. 02-0200. A duplicate copy of this paper is enclosed.

I, Richard E. Fichter, hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine, or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the above-captioned patent.

Respectfully submitted,

BACON & THOMAS, PLLC

By:______ Richard E. Fichter Registration No. 26,382

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REF:kdd Terminal Disclaimer.wpd

January 4, 2005

ANHANG ZUM RECHERCHENBERICHT ÜBER DIE STANDARDRECHERCHE NR.

RS 111292 DE 10344229

In diesem Anhang sind die Mitglieder der Patentfamilien der im obengenannten Recherchenbericht angeführten Patentdokumente angegeben.
Die Angaben über die Familienmitglieder entsprechen dem Stand der Datei des Europäischen Patentamts am Diese Angaben dienen nur zur Unterrichtung und erfolgen ohne Gewähr.

13-07-2004

im Rechorchenbericht angeführtes Patentdokument			Datum der Veröffentlichung		Mitglied(er) der Patentamille	Datum der Veröffentlichur
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EPO FORM PONT

Für nähere Einzelheiten zu diesem Anhang : siehe Amtsblatt des Europäischen Patentante, Nr.12/82

2

In diesem Bescheid sind folgende Entgegenhaltungen erstmalig genannt. (Bei deren Nummerierung gilt diese auch für das weitere Verfahren):

- 1) EP 0 329 120 A2 2) WO 86/06 488 A1 ν 3) EP 0 583 833 A2 4) DE 101 37 565 A1 5) DE 43 41 862 A1 > 6) US 5 362 654 7) US 5 147 780 8) US 38 25 410 4 9) US 4 324 758 10) US 3 888 629 11) WO 99/01221 A1 12) US 3 704 206 🗸 13) DE 195 45 130 A1 14) US 4 526 690 » 15) US 5 364 591 🛩 16) US 5 863 502 🗸
- 1. Aus der Druckschrift 1) (Figur 3 samt Beschreibung) ist eine Vorrichtung mit allen Merkmalen des Anspruchs 1 bekannt, der Anspruch 1 ist aus diesem Grunde wegen mangelnder Neuheit seines Gegenstandes nicht gewährbar.
- 2. Ähnliche Vorrichtungen sind auch aus den Druckschrift 2) bis 10) bekannt.
- Aus der Druckschrift 1) ist auch ein Verfahren gem

 äß Anspruch 22 bekannt, der Anspruch 22 ist somit wegen mangelnder Neuheit seines Gegenstandes ebenfalls nicht gewährbar.
- 4. Bezüglich der Merkmale der Unteransprüche 2 bis 21 wird neben den Druckschriften 1) bis 10) noch auf die Druckschriften 11) bis 16) verwiesen.
- 5. Es ist nicht ersichtlich, dass durch die Aufnahme von Merkmalen der Unteransprüche in

Process and apparatus for the determination of parameters of a breath condensate

Veröffentlichungsnummer DE10137565

Auch veröffentlicht als

US2003023389 (A

Erfinder

Veröffentlichungsdatum: 2003-03-06

DIETZE STEFAN (DE); ROTHE MICHAEL (DE);

BECHER GUNTHER (DE); LEHMANN CLAUDIA

(DE); PORZELT JOACHIM (DE)

Anmelder:

FILT LUNGEN UND THORAXDIAGNOSTI (DE)

Klassifikation:

- Internationale:

G01N33/497; A61B5/083

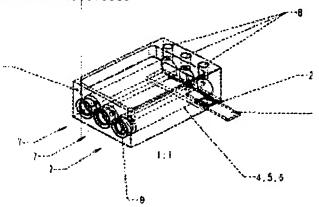
- Europäische:

G01N33/497

Anmeldenummer: Prioritätsnummer(n): DE20011037565 20010730 DE20011037565 20010730

Keine Zusammenfassung verfügbar für DE10137565 Zusammenfassung der korrespondierenden Patentschrift US2003023389

The invention relates to a process and apparatus for the determination of the parameters of a breath condensate using at least one sensor for the measurement of the parameters and an analysis unit with display for the measurement results. The at least one sensor (2) is positioned in a closed cassette (1), whereby the cassette includes storage containers (4, 5, 6) with flushing solutions and/or solutions for the conditioning of the at least one sensor and/or calibration solutions and/or compounds for the manufacture of a calibration solution and/or for the dilution of the sample solution and/or for elevation of the ion concentration or the conductivity of the sample solution. The storage containers (4, 5, 6) deliver their contents completely or in metered amounts into the cassette and/or onto the at least one sensor (2) under the action of means acting from the outside and through the cassette walls. Furthermore, the cassette (1) includes at least one opening for the introduction of the sample solution onto the at least one sensor (2), and has contacts (3) for receiving the measurement data originating from the at least one sensor (2)



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Auch veröffentlicht als

US5500187 (A

JP7191032 (A)

Sample assay device in card form - useful for cocaine detection by latex agglutination test

Veröffentlichungsnummer DE4341862

Veröffentilchungsdatum: 1994-06-09

Erfinder

JOHNSON SCOTT DAVID (US); KAPRAUN RICHARD CHARLES (US); LACOUR KEYIN EUGENE (US); RODRIGUEZ II NOE ESAU (US); DEOMS JAMES HOWARD (US); LEGINUS JOSEPH M (US); MILEAF DARYL SUNNY (US);

YOUNG RICHARD MCLEOD (US)

Anmeider:

WESTINGHOUSE ELECTRIC CORP (US)

Klassifikation: - Internationale:

G01N35/00; G01N33/94; G01N33/577

- Europăische: G01N33/543D; B01L3/00C6; G01N33/543K;

G01N35/00C

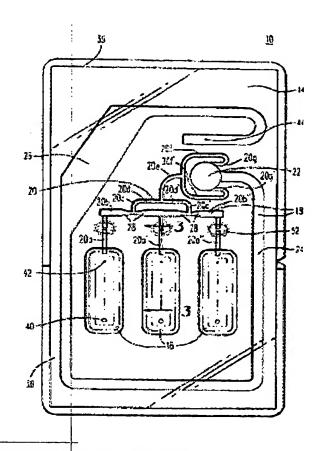
Anmeldenummer: Prioritätsnummer(n):

DE19934341862 19931208 US19920986816 19921208

Zusammenfassung von DE4341862

An assay device, for analysing a sample of an unknown substance by mixing with a reagent, has (a) a housing with one or more reagent holding reservoirs; (b) an inlet opening for accommodating the sample in the housing; and (c) at least one path which is used to control reagent flow rate and mixing characteristics and which produces a fluid connection between the reagent and the inlet opening for mixing with the sample.

A method of determining the identity of an unknown sample involves (i) providing an assay card comprising the above assay device including a reaction path with an optical viewing face for observing the result of mixing reagent and sample; (ii) introducing the sample into the assay device; (iii) pressing reagent out of the reservoir through the control path into the sample-contg, part of the housing at a first flow rate; and (iv) reducing the flow rate so that the reagent and sample mixt. flows from the samplecontg. part of the housing into the reaction path. USE/ADVANTAGE - E.g. for latex agglutination reactions using monoclonal mouse anti-cocaine antibodies in buffered soln, as the antibody reagent and latex-cocaine particles in buffered soln. as the latex reagent, for cocaine detection. The device allows detection and measurement of an analyte in a sample or on a surface and is an inexpensive throw-away unit which facilitates sample taking and reaction by the user.



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http://v3.espacenet.com/textdoc?DB=EPODOC&IDX=DE4341862&F=0&QPN=DE4... 22.12.2004

Auch veröffentlicht als

WO9721095 (A:

WO9721095 (A:

Verfahren und Vorrichtungen für ein modulares Mikrosystem für hochgenauchemische Schnell-Analysen und Verfahren zur Herstellung

Veröffentlichungsnummer DE19545130

Veröffentlichungsdatum: 1997-06-05

Erfinder

CAMMANN KARL PROF DR (DE)

Anmelder:

CAMMANN KARL PROF DR (DE)

Klassifikation:

- Internationale:

G01N31/20; G01N31/16; G01N31/22; G01N33/53; G01N27/416; B01L3/02; G01N33/50; C12Q1/00;

C12Q1/26; C12Q1/32; C12Q1/42; C12Q1/54

- Europäische:

B01L3/00C2D; G01N21/29B; G01N31/16

Anmeldenummer: Prioritätsnummer(n):

DE19951045130 19951204 DE19951045130 19951204

Zusammenfassung von DE19545130

The invention relates to a method and a device for determining the concentration of an analyte, analyte mixture or biochemical analyte-equivalent substance (titrant) contained in a sample using at least one selective reagent (titrator). Instead of the concentration-proportional intensity measurement known from prior art, the invention uses an absolute equivalent quantity measurement which can be miniaturised and can be read off with considerably greater accuracy. Measurement of the titrator quantity equivalent to the analyte concentration, i.e. of the chemical titrator equivalent to be measured, is reduced to a simple and rapid procedure for determining a location or line. The reaction in the reaction series, i.e. the point on a titration line, at which equivalent quantities of analyte and titrator are present is determined. Other process steps such as biochemical analyte-identification reactions or chromatographic separations can also be included.

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